

Result No.	Score	Query Match	Length	DB ID	Description
1	49	100.0	9	4 AAB1114	Aab1114 Human MUC
2	49	100.0	9	5 ABG79089	Abg79089 Human MUC
3	49	100.0	9	6 ADA50588	Ada50588 Mucin 1 (
4	49	100.0	9	8 ADG89655	Adg89655 Class I H
5	49	100.0	9	8 ADG20359	Adg20359 Antigenic
6	49	100.0	13	2 AAW77232	Aaw77232 Peptide 8
7	49	100.0	20	8 ADEA3990	Ade43990 MUC-1 imp
8	49	100.0	20	8 ADP22621	Adf22621 MUC-1 imp
9	49	100.0	30	5 AAU84987	Aau84987 Human MUC
10	49	100.0	173	3 AAY71021	Aay71021 Human MUC
11	49	100.0	180	2 AAR27664	Aar27664 C-terminal
12	49	100.0	256	8 AD157759	Adi57759 Human bre
13	49	100.0	287	2 AAR27665	Aar27665 Secreted
14	49	100.0	295	3 AAY71027	Aay71027 Ubiquitin
15	49	100.0	307	6 ADA50571	Ada50571 Mucin 1 (
16	49	100.0	312	5 AAU84810	Aau84810 Human MUC
17	49	100.0	316	8 AD157755	Adi57755 Human bre
18	49	100.0	325	8 AD157777	Adi57777 Human bre
19	49	100.0	327	2 AAR6298	Aar6298 Glycoprot
20	49	100.0	336	8 ADI57782	Adi57782 Human bre
21	49	100.0	348	2 AAR27662	Aar27662 C-terminal
22	49	100.0	350	8 ADI57754	Adi57754 Human bre
23	49	100.0	370	8 ADI57758	Adi57758 Human bre
24	49	100.0	379	8 ADI57779	Adi57779 Human bre
25	49	100.0	396	8 ADI57776	Adi57776 Human bre

ALIGNMENTS

RESULT 1
AAB1114
ID AAB1114 standard; peptide; 9 AA.

AAB1114;
XX
AC
XX
DT 16-FEB-2001 (first entry)
XX
Human MUC-1 protein fragment SEQ ID NO 1.
DE
XX
KW Human; MUC-1; tumor; HLA-A2 restricted; immune reaction; treatment;
XX
human leukocyte antigen; gene therapy; antigen-presenting cell.
OS Homo sapiens.
XX
PN DE19917195-A1.
XX
PD 19-OCT-2000.
XX
PP 16-APR-1999;
XX
PR 16-APR-1999;
XX
XX (UYTU-) UNIV TUEBINGEN EBERHARD-KARLS.

PA
XX
PI
XX
DR 2001-03-2872/05.
XX
PT New peptide derived from the MUC-1 tumor marker, used to induce a
cytotoxic T cell response for treatment or prevention of tumors.
XX
PP 16-APR-1999;
XX
PR 16-APR-1999;
XX
XX (UYTU-) UNIV TUEBINGEN EBERHARD-KARLS.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	49	100.0	9	4 AAB1114	Aab1114 Human MUC
2	49	100.0	9	5 ABG79089	Abg79089 Human MUC
3	49	100.0	9	6 ADA50588	Ada50588 Mucin 1 (
4	49	100.0	9	8 ADG89655	Adg89655 Class I H
5	49	100.0	9	8 ADG20359	Adg20359 Antigenic
6	49	100.0	13	2 AAW77232	Aaw77232 Peptide 8
7	49	100.0	20	8 ADEA3990	Ade43990 MUC-1 imp
8	49	100.0	20	8 ADP22621	Adf22621 MUC-1 imp
9	49	100.0	30	5 AAU84987	Aau84987 Human MUC
10	49	100.0	173	3 AAY71021	Aay71021 Human MUC
11	49	100.0	180	2 AAR27664	Aar27664 C-terminal
12	49	100.0	256	8 AD157759	Adi57759 Human bre
13	49	100.0	287	2 AAR27665	Aar27665 Secreted
14	49	100.0	295	3 AAY71027	Aay71027 Ubiquitin
15	49	100.0	307	6 ADA50571	Ada50571 Mucin 1 (
16	49	100.0	312	5 AAU84810	Aau84810 Human MUC
17	49	100.0	316	8 AD157755	Adi57755 Human bre
18	49	100.0	325	8 AD157777	Adi57777 Human bre
19	49	100.0	327	2 AAR6298	Aar6298 Glycoprot
20	49	100.0	336	8 ADI57782	Adi57782 Human bre
21	49	100.0	348	2 AAR27662	Aar27662 C-terminal
22	49	100.0	350	8 ADI57754	Adi57754 Human bre
23	49	100.0	370	8 ADI57758	Adi57758 Human bre
24	49	100.0	379	8 ADI57779	Adi57779 Human bre
25	49	100.0	396	8 ADI57776	Adi57776 Human bre

This invention describes a novel peptide (I) derived from the MUC-1 gene which is able to induce an immune response against tumor cells (II) or the nucleic acid (II) encoding (I), used to induce an immune response against tumor cells, so are useful for treatment or prevention of tumors, in conjunction with other tumor therapies. In particular (II) is used in gene therapy or for in vitro transfection or transformation of cells (particularly antigen-presenting cells, optionally in vivo), for expression of (I). (I) has a high binding capacity for HLA-A2 and can reverse the usual suppression of the immune response associated with tumor cells. By introducing the nucleic acid that encodes (I) into an antigen-presenting cell in vitro, then returning the cells to the patient, a more certain and controlled response is achieved, compared with administration of the peptide plus adjuvant

XX Sequence 9 AA;
 SQ Query Match 100.0%; Score 49; DB 4; Length 9;
 Best Local Similarity 100.0%; Pred. No. 1.7e+06;
 Matches 9; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 Qy 1 STAPPVHNV 9
 Db 1 STAPEVHNV 9

RESULT 2
 ID ABG79089 standard; peptide; 9 AA.
 AC ABG79089;
 XX DT 15-NOV-2002 (first entry)
 DE Human MUC1 Class I HLA widely expressed antigen peptide #2.
 XX Cell penetrating peptide; cancer; tumour; melanoma; thymoma; antigen;
 KW lymphoma; sarcoma; lung cancer; non-Hodgkin's lymphoma; leukemia;
 KW Hodgkin's lymphoma; uterine cancer; cervical cancer; bladder cancer;
 KW kidney cancer; adenocarcinoma; breast cancer; prostate cancer;
 KW ovarian cancer; pancreatic cancer; epitope; vaccine; dendritic cell;
 KW tumour infiltrating lymphocyte; TIL; human leukocyte antigen; HLA-1;
 KW cytostatic; human.
 XX OS Homo sapiens.
 PN WO20026057-A2.
 XX PD 22-AUG-2002.
 XX PP 15-FEB-2002; 2002WO-US005212.
 PR 15-FEB-2001; 2001US-0268687P.
 PA (BAYU) BAYLOR COLLEGE MEDICINE.
 XX PI Wang R;
 XX DR WPI; 2002-627577/67.
 XX PR Novel composition for treating a disease in an animal, comprises an
 PR immune effector cell and cell penetrating peptide associated with an
 PR antigen or antibody.
 XX Disclosure; Page 18; 61pp; English.
 XX The invention relates to a composition (I) comprising an immune effector
 cell and a cell penetrating peptide (CPP) associated with an antigen or
 antibody. Also included are (1) a vaccine comprising (I), CPP associated
 with an antigen, and a pharmaceutically acceptable carrier and (2)
 preparing a composition for a disease, by providing (I) and CPP
 associated with an antigen for disease, and introducing the antigen-
 associated CPP to (I), where antigen enters into the cell. The antigens
 are, for example, tumour antigen derived epitopes recognised by tumour
 infiltrating lymphocytes (TIL) of HLA (human leukocyte antigen) class I
 or II. The composition is useful for enhancing immunity in an animal to a
 disease, by administering a mature dendritic cell comprising CPP
 associated with an antigen to disease, to the animal, such that following
 the administration, animal is protected from disease, where the animal
 comprises both CD4+ and CD8+ T cells. It is also useful for treating a
 disease (e.g. cancer, tumour, melanoma, thymoma, sarcoma, lung
 cancer, non-Hodgkin's lymphoma, Hodgkin's lymphoma, uterine
 cancer, cervical cancer, bladder cancer, kidney cancer, adenocarcinoma,
 breast cancer, prostate cancer, ovarian cancer and pancreatic cancer).
 The animal is further subjected to a cancer treatment including surgery,
 radiation, chemotherapy or gene therapy. The administration of (I),
 preferably dendritic cell is prior to, subsequent to or concurrent with,

CC the cancer treatment. The present sequence is a tumour antigen derived
 CC epitope for inclusion in the composition of the invention.
 XX SQ Sequence 9 AA;
 Query Match 100.0%; Score 49; DB 5; Length 9;
 Best Local Similarity 100.0%; Pred. No. 1.7e+06;
 Matches 9; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 Qy 1 STAPPVHNV 9
 Db 1 STAPPVHNV 9

RESULT 3
 ID ADA50588
 XX AC ADA50588;
 XX DT 20-NOV-2003 (first entry)
 XX DE Mucin 1 (MUC-1) CTL epitope, SEQ ID NO:43.
 KW Nucleic acid vaccine; DNA vaccine; tumour antigen; cytokine adjuvant;
 KW humoral response; cellular response; immune response; immunotherapy;
 KW cancer; cytotoxic; vaccine; gene therapy; mucin 1; MUC-1;
 KW cytotoxic T lymphocyte; CTL epitope.
 XX OS Unidentified.
 PN WO2003031569-A2.
 XX PD 17-APR-2003.
 XX PP 18-SEP-2002; 2002WO-US029640.
 XX PR 10-OCT-2001; 2001US-0328371P.
 XX PA (CENZ) CENTOCOR INC.
 XX PI Snyder L, Scallan B, Knight DM, McCarthy SG, Goletz TJ;
 PI Branigan PJ;
 XX DR WPI; 2003-393437/37.
 XX PT New nucleic acid vaccine, useful for eliciting an immune response to a
 PT cancer associated tumor protein in a mammal.
 XX PS Claim 1a; Page 45; 92pp; English.
 XX The invention relates to a nucleic acid vaccine comprising one or more
 CC tumour antigen-encoding nucleic acids and one or more cytokine adjuvant-
 CC encoding nucleic acids. The tumour antigen encoded by the vaccine is
 CC mucin 1 (MUC-1), the kallikrein KLK2, or prostate specific antigen (PSA,
 CC also known as Klk3), and the cytokine adjuvant encoded can be interleukin
 CC -12 (IL-12), granulocyte macrophage-colony stimulating factor (GM-CSF),
 CC or especially interleukin-18 (IL-18). The antigen-encoding nucleic acid
 CC is preferably under the control of a promoter such as the cytomegalovirus
 CC immediate early promoter, the dihydrofolate reductase promoter or the
 CC early or late SV40 promoters. The invention also encompasses the method
 CC of eliciting an immune response to a tumour antigen in a mammal using the
 CC vaccine of the invention. Coexpression of the antigen and adjuvant
 CC induces a humoral or cellular response to the tumour antigen generating
 CC an immune response useful for treatment or prophylaxis of cancers. The
 CC present sequence represents a mucin 1 (MUC-1) polypeptide sequence which
 CC is specifically claimed for use in the vaccine of the invention.
 XX SQ Sequence 9 AA;
 Query Match 100.0%; Score 49; DB 6; Length 9;
 Best Local Similarity 100.0%; Pred. No. 1.7e+06;
 Matches 9; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 STAPPVHN 9
Db 1 STAPPVHN 9

RESULT 4
ID ADG89655
ID ADG89655 standard; peptide; 9 AA.

AC ADG89655;

DT 11-MAR-2004 (first entry)

DB Class I HLA-restricted widely expressed antigen #20.

XX metastatic cancer cell differentiation; mutated fibronectin; RW metastatic cancer; class I HLA-restricted; widely antigen.

Unidentified.

W2003100027-A2
PN W2003100027-A2

XX PD-DEC-2003.

XX PF 28-MAY-2003; 2003WO-US016736.

XX PR 28-MAY-2002; 2002US-0383530P.

PA (BAYU) BAYLOR COLLEGE MEDICINE.

XX PI Wang R;

XX DR WIT; 2004-035134/03.

XX Identifying a cell that differentiates into a metastatic cancer cell, PR useful for preventing metastatic cancer, comprises identifying a mutated PR fibronectin in the cell.

XX Disclosure; SEQ ID NO 98; 137pp; English.

XX The invention comprises a method for identifying a cell that will differentiate into a metastatic cancer cell, the method involves identifying a mutated fibronectin in the cell. The method of the invention is useful for preventing metastatic cancer. The present amino acid sequence represents a Class I HLA-restricted widely expressed

XX Sequence 9 AA;

Query Match 100.0%; Score 49; DB 8; Length 9;
Best Local Similarity 100.0%; Pred. No. 1.7e+06;
Matches 9; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 STAPPVHN 9
Db 1 STAPPVHN 9

RESULT 5
ID ADG20359
ID ADG20359 standard; peptide; 9 AA.

AC ADG20359;

DT 11-MAR-2004 (first entry)

DB Antigenic peptide SEQ ID NO:35.

XX double-chimeric beta 2-microglobulin; antigenic peptide;
XX antigen-presenting cell; beta 2-microglobulin;
XX major histocompatibility complex class I epitope; MHC class I epitope;
XX cytosatic; antibacterial; complex class I epitope; MHC class I epitope;
XX protozoaide; vaccine;

KW cytotoxic T lymphocyte induction; cancer; pathogenic organism;
XX tumour associated antigen; pathogenic antigen.
OS Synthetic.

XX PN WO2003106616-A2.

XX PD 24-DEC-2003.

XX PF 12-JUN-2003; 2003WO-TL000501.

XX PR 12-JUN-2002; 2002US-0388233P.

XX PA (GAVI-) GAVISH-GALILEE BIO APPL LTD.

XX Gross G, Margalit A;

XX DR 2004-071554/07.

XX Novel double-chimeric beta2-microglobulin polynucleotide useful for PT treating cancer, comprising sequence encoding polypeptide capable of PT presentation of antigenic peptides.

XX PS Claim 16; SEQ ID NO 35; 86pp; English.

XX The present invention describes a double-chimeric beta 2-microglobulin polynucleotide (I) comprising a polypeptide (II) that is capable of high level presentation of antigenic peptides on antigen-presenting cells, where (II) comprising a beta 2-microglobulin molecule that is linked through its carboxyl terminal to a polypeptide stretch which allows the anchorage of the beta 2-microglobulin molecule to the cell membrane, and through its amino terminal to a antigenic peptide comprising major histocompatibility complex (MHC) class I epitope. The antigenic peptide is not related to an autoimmune disease. Also described: (1) an expression vector (III) comprising (I) and is a recombinant viral vector; (2) an antigen-presenting cell (IV) transfected with (I); (3) a DNA vaccine (V) comprising a (I) or (III) or (IV) vaccine (VI) for the prevention or treatment of cancer comprising (IV) which express (I) or tumour cells transfected with (I), where the cells have been pulsed with an antigenic peptide derived from one tumour associated antigen; and (5) a pharmaceutical composition (VII) comprising (I), (III) or (IV) as an active ingredient and carrier. (I) has cytosatic, antibacterial, virucide, fungicide and protozoaide activities, and can be used in vaccines, and for inducing cytotoxic T lymphocytes. (I) and (V) can be used for the prevention or treatment of cancer or for a disease caused by a pathogenic organism. (VI) is useful for prevention or treatment of cancer, or disease caused by a pathogenic organism, where (VI) presents one tumour associated antigen, or tumour associated antigen. (VI) is also useful for immunising a mammal against a pathogenic organism, which involves immunising the mammal with (VI). (I) is useful for inducing class I-restricted CTL response in a mammal. The present sequence is used in the exemplification of the present invention.

XX SQ Sequence 9 AA;

Query Match 100.0%; Score 49; DB 8; Length 9;
Best Local Similarity 100.0%; Pred. No. 1.7e+06;
Matches 9; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 STAPPVHN 9
Db 1 STAPPVHN 9

RESULT 6
ID AAW77232
ID AAW77232 standard; peptide; 13 AA.

XX AC AAW77232;

XX DT 20-NOV-1998 (first entry)

DE Peptide sequence encoding MUC1 tandem repeat unit c.
 XX MUC1; recombinant pox virus; cytotoxic T-lymphocyte; immunogen; tumour;
 KW tumour-associated antigen.
 XX Homo sapiens.
 XX WO9837095-A2.
 XX PD 27-AUG-1998.
 XX DF 24-FEB-1998; 9BW0-US003693.
 XX PR 24-FEB-1997; 97US-0038253P.
 XX PA (THERION BIOLOGICS CORP.
 PA (USSH) US DEPT HEALTH & HUMAN SERVICES.
 PA (DAND) DANA FARBER CANCER INST INC.
 XX P1 Schliom J, Kantor J, Kufe D, Panicali D, Gritz L;
 XX DR; 1998-467492/40.
 XX PT New recombinant pox virus for tumour therapy - comprises DNA encoding an
 PR immunogenic mini-MUC1 fragment comprising 5-25 MUC1 tandem repeat units.
 XX Example 1; Page 20; 42PP; English.
 XX The MUC1 tandem repeat units AAW77230-W77232 were used to create an
 CC immunogenic mini-MUC1 fragment for inclusion in a recombinant pox virus
 CC (RPV). The RPV was used in a pharmaceutical composition also containing
 CC an immunomodulator to generate MUC1 specific cytotoxic T-lymphocytes. The
 CC recombinant pox virus therefore encodes an immunogenic MUC1 fragment that
 CC does not undergo significant genetic deletion, thereby providing an
 CC unexpectedly stable and immunogenic pox virus. They can be used to
 CC prevent or treat tumours expressing MUC1 tumour-associated antigens
 XX Sequence 13 AA;
 Query Match 100.0%; Score 49; DB 2; Length 13;
 Best Local Similarity 100.0%; Pred. No. 0.14; Indels 0; Gaps 0;
 Matches 9; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 SQ

XX Burden N, Hamblin P;
 XX WPI; 2004-035056/03.
 XX PT New nucleic acid molecule encoding a MUC-1 derivative that is devoid of
 PT all perfect repeats, useful as vaccine for treating or preventing MUC-1
 PT expressing tumors e.g. carcinoma of the breast, lung or gastrointestinal
 PT carcinomas.
 XX Example; Page 16; 34PP; English.
 XX PS
 XX CC The present invention describes a nucleic acid molecule encoding a MUC-1
 CC derivative that is devoid of all perfect repeats. Also described is:
 CC a plasmid comprising the DNA molecule; (2) a protein encoded by a nucleic
 CC acid molecule; (3) a pharmaceutical composition comprising the nucleic
 CC acid, the plasmid or the protein and a pharmaceutical acceptable
 CC excipient, diluent or carrier; and (4) a method of treating or preventing
 CC tumours. MUC-1 has cytotoxic activity, and can be used in vaccines. The
 CC nucleic acid, plasmid, a protein or the pharmaceutical composition of the
 CC present invention can be used in medicine. The nucleic acid or the
 CC protein can be used in the preparation of a medicament for the treatment
 CC or prevention MUC-1 expressing tumours. The tumour can be carcinomas of
 CC the breast, lung, Gastric or other gastrointestinal carcinomas. The
 CC nucleic acid vaccines are easy to produce in large quantities compared
 CC over conventional protein vaccination. Even at small doses they have been
 CC reported to induce strong immune responses and can induce a cytotoxic T
 CC lymphocyte immune response as well as an antibody response. The present
 CC sequence represents a MUC-1 imperfect repeat peptide, which is used in
 CC the exemplification of the present invention.
 XX Sequence 20 AA;
 Query Match 100.0%; Score 49; DB 8; Length 20;
 Best Local Similarity 100.0%; Pred. No. 0.21; Indels 0; Gaps 0;
 Matches 9; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 SQ

XX Query 1 STAPPVHN 9
 XX Database ADF32621 standard; peptide; 20 AA.
 XX ID ADF32621;
 XX AC ADF32621;
 XX DT 26-FEB-2004 (first entry)
 XX DE MUC-1 imperfect repeat 4 VNTR.
 XX KW MUC-1 antigen; immune response; MUC-1; variable number of tandem repeat;
 KW VNTR; repeat unit; tumour; metastasis; cytostatic; vaccine; gene therapy.
 XX OS Synthetic.
 XX PN WO2003100060-A2.
 XX PR 24-MAY-2002; 2002GB-00012046.
 XX PD 04-DEC-2003.
 XX PA (GLAXO GROUP LTD.
 XX PI Burden N, Ellis JH, Hamblin PA;
 XX DR WPI; 2004-042811/04.
 XX PT New nucleic acid molecule encoding a MUC-1 antigen, useful for preparing
 PT a composition for treating or preventing tumors or metastases.
 PA (GLAXO GROUP LTD.

XX Burden N, Hamblin P;
 XX WPI; 2004-035056/03.
 XX PT New nucleic acid molecule encoding a MUC-1 derivative that is devoid of
 PT all perfect repeats, useful as vaccine for treating or preventing MUC-1
 PT expressing tumors e.g. carcinoma of the breast, lung or gastrointestinal
 PT carcinomas.
 XX Example; Page 16; 34PP; English.
 XX PS
 XX CC The present invention describes a nucleic acid molecule encoding a MUC-1
 CC derivative that is devoid of all perfect repeats. Also described is:
 CC a plasmid comprising the DNA molecule; (2) a protein encoded by a nucleic
 CC acid molecule; (3) a pharmaceutical composition comprising the nucleic
 CC acid, the plasmid or the protein and a pharmaceutical acceptable
 CC excipient, diluent or carrier; and (4) a method of treating or preventing
 CC tumours. MUC-1 has cytotoxic activity, and can be used in vaccines. The
 CC nucleic acid, plasmid, a protein or the pharmaceutical composition of the
 CC present invention can be used in medicine. The nucleic acid or the
 CC protein can be used in the preparation of a medicament for the treatment
 CC or prevention MUC-1 expressing tumours. The tumour can be carcinomas of
 CC the breast, lung, Gastric or other gastrointestinal carcinomas. The
 CC nucleic acid vaccines are easy to produce in large quantities compared
 CC over conventional protein vaccination. Even at small doses they have been
 CC reported to induce strong immune responses and can induce a cytotoxic T
 CC lymphocyte immune response as well as an antibody response. The present
 CC sequence represents a MUC-1 imperfect repeat peptide, which is used in
 CC the exemplification of the present invention.
 XX Sequence 20 AA;
 Query Match 100.0%; Score 49; DB 8; Length 20;
 Best Local Similarity 100.0%; Pred. No. 0.21; Indels 0; Gaps 0;
 Matches 9; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 SQ

XX Query 1 STAPPVHN 9
 XX Database ADF32621 standard; peptide; 20 AA.
 XX ID ADF32621;
 XX AC ADF32621;
 XX DT 26-FEB-2004 (first entry)
 XX DE MUC-1 imperfect repeat 4 VNTR.
 XX KW MUC-1 antigen; immune response; MUC-1; variable number of tandem repeat;
 KW VNTR; repeat unit; tumour; metastasis; cytostatic; vaccine; gene therapy.
 XX OS Synthetic.
 XX PN WO2003100060-A2.
 XX PR 24-MAY-2002; 2002GB-00012046.
 XX PD 04-DEC-2003.
 XX PA (GLAXO GROUP LTD.
 XX PI Burden N, Ellis JH, Hamblin PA;
 XX DR WPI; 2004-042811/04.
 XX PT New nucleic acid molecule encoding a MUC-1 antigen, useful for preparing
 PT a composition for treating or preventing tumors or metastases.
 PA (GLAXO GROUP LTD.

XX Burden N, Hamblin P;
 XX WPI; 2004-035056/03.
 XX PT New nucleic acid molecule encoding a MUC-1 derivative that is devoid of
 PT all perfect repeats, useful as vaccine for treating or preventing MUC-1
 PT expressing tumors e.g. carcinoma of the breast, lung or gastrointestinal
 PT carcinomas.
 XX Example; Page 16; 34PP; English.
 XX PS
 XX CC The present invention describes a nucleic acid molecule encoding a MUC-1
 CC derivative that is devoid of all perfect repeats. Also described is:
 CC a plasmid comprising the DNA molecule; (2) a protein encoded by a nucleic
 CC acid molecule; (3) a pharmaceutical composition comprising the nucleic
 CC acid, the plasmid or the protein and a pharmaceutical acceptable
 CC excipient, diluent or carrier; and (4) a method of treating or preventing
 CC tumours. MUC-1 has cytotoxic activity, and can be used in vaccines. The
 CC nucleic acid, plasmid, a protein or the pharmaceutical composition of the
 CC present invention can be used in medicine. The nucleic acid or the
 CC protein can be used in the preparation of a medicament for the treatment
 CC or prevention MUC-1 expressing tumours. The tumour can be carcinomas of
 CC the breast, lung, Gastric or other gastrointestinal carcinomas. The
 CC nucleic acid vaccines are easy to produce in large quantities compared
 CC over conventional protein vaccination. Even at small doses they have been
 CC reported to induce strong immune responses and can induce a cytotoxic T
 CC lymphocyte immune response as well as an antibody response. The present
 CC sequence represents a MUC-1 imperfect repeat peptide, which is used in
 CC the exemplification of the present invention.
 XX Sequence 20 AA;
 Query Match 100.0%; Score 49; DB 8; Length 20;
 Best Local Similarity 100.0%; Pred. No. 0.21; Indels 0; Gaps 0;
 Matches 9; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 SQ

XX Query 1 STAPPVHN 9
 XX Database ADF32621 standard; peptide; 20 AA.
 XX ID ADF32621;
 XX AC ADF32621;
 XX DT 26-FEB-2004 (first entry)
 XX DE MUC-1 imperfect repeat 4 VNTR.
 XX KW MUC-1 antigen; immune response; MUC-1; variable number of tandem repeat;
 KW VNTR; repeat unit; tumour; metastasis; cytostatic; vaccine; gene therapy.
 XX OS Synthetic.
 XX PN WO2003100060-A2.
 XX PR 24-MAY-2002; 2002GB-00012046.
 XX PD 04-DEC-2003.
 XX PA (GLAXO GROUP LTD.
 XX PI Burden N, Ellis JH, Hamblin PA;
 XX DR WPI; 2004-042811/04.
 XX PT New nucleic acid molecule encoding a MUC-1 antigen, useful for preparing
 PT a composition for treating or preventing tumors or metastases.
 PA (GLAXO GROUP LTD.

XX Disclosure; Page 2; 66pp; English.

XX The present invention describes a nucleic acid molecule which encodes a MUC-1 antigen. The nucleic acid is capable of raising an immune response in vivo, has reduced susceptibility to recombination than full-length MUC -1 and comprises between 1 and 15 variable number of tandem repeats (VNTR) perfect repeat units. Also described: (1) a plasmid comprising the DNA molecule; (2) a protein encoded by the nucleic acid; (3) a pharmaceutical composition comprising the nucleic acid, plasmid or protein and an excipient, diluent or carrier; and (4) a method of treating or preventing tumours or metastases. A MUC1 antigen has cytostatic activity, and can be used in vaccines, and in gene therapy. The nucleic acid is useful for preparing a composition for treating or preventing tumours or metastases. The present sequence is used in the exemplification of the present invention.

XX Sequence 20 AA;

Query	Match	Score	DB	Length
Qy	1 STAPPVNV 9	100.0%	8	20;
Db	10 STAPPVNV 18	100.0%	8	20;

XX Sequence 20 AA;

Query	Match	Score	DB	Length
Qy	1 STAPPVNV 9	100.0%	5	30;
Db	9 STAPPVNV 17	100.0%	5	30;

XX Sequence 30 AA;

Query	Match	Score	DB	Length
Qy	1 STAPPVNV 9	100.0%	5	30;
Db	9 STAPPVNV 17	100.0%	5	30;

XX Sequence 30 AA;

RESULT 9

AAU84987 ID AAU84987 Standard; peptide; 30 AA.

XX AC AAU84987;

XX DT 08-MAY-2002 (first entry)

XX DE Human MUC1R segment 1.

XX KW Savine vaccine; cancer; viral infection; HIV; hepatitis C virus; viral infection; human immunodeficiency virus; melanoma; bacterial infection; Salmonella; Legionella; parasitic infection; Trypanosoma; Toxoplasma; Giardia.

XX OS Homo sapiens.

XX PN WO20025827-A2.

XX PD 11-MAY-2000.

XX PP 18-OCT-1999;

XX PR 30-OCT-1998;

XX PA (MENA) MENARINI RICERCHE SPA.

XX PI Parente D, Di Massimo AM, De Santis R;

XX DR WPI; 2000-365410/31.

XX DR N-PSDB; AAD00385.

XX PT Composition containing one or more DNA molecules encoding fragments of a

PT tumor therapy.

XX PS Claim 16; Fig 2; 56pp; English.

XX The present sequence is a fragment of human Mucin 1 (MUC-1), an antigenic

CC protein overexpressed in tumor cells. The sequence was obtained from

CC BT20 tumor cells. The corresponding DNA sequence is cloned into a pMRS30

CC expression vector and used in pharmaceutical composition e.g. vaccine for

CC inducing an antigen-specific anti-tumour immune response. Composition

CC containing this DNA molecule is useful in anti-tumour therapy. The

CC function associated with the parent polypeptide and for inducing an

CC immune response against a pathogen or cancer. Also included are a

XX

SQ Sequence 173 AA;
 Query Match 100.0%; Score 49; DB 3; Length 173;
 Best Local Similarity 100.0%; Pred. No. 1.9;
 Matches 9; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 AC AAR27664;
 XX DT 25-MAR-2003 (revised)
 AC DT 06-NOV-1992 (first entry)
 XX DE C-terminal region of H23-ETA-S antigen.
 XX KW Secreted; human epithelial antigen; Monoclonal antibody H23; vaccine;
 XX malignant tumour; breast cancer; tandem repeat.
 OS Homo sapiens.
 XX Key Location/qualifiers
 FT 1..40 /note= "contains 2 tandem repeats - can have up to 80
 copies"
 FT Misc-difference 7
 FT /label= Pro, Ala
 FT /note= "natural polymorphism"
 FT Misc-difference 17
 FT /label= Thr, Asn
 FT /note= "natural polymorphism"
 FT Misc-difference 20
 FT /label= Pro, Ala
 FT /note= "natural polymorphism"
 FT Misc-difference 27
 FT /label= Pro, Ala
 FT /note= "natural polymorphism"
 FT Misc-difference 37
 FT /label= Pro, Ala
 FT /note= "natural polymorphism"
 FT Misc-difference 40
 FT /label= Thr, Asn
 FT /note= "natural polymorphism"
 XX WO9207000-A1.
 XX PD 30-APR-1992.
 XX PP 23-OCT-1991; 91WO-FR0000835.
 XX PR 23-OCT-1990; 90FR-00013101.
 XX PN (TRGE) TRANSGENE SA.
 XX PA Chambon P, Kiery MP, Lathe R, Hareveni M;
 XX WPI; 1992-167097/20.
 XX DR N-PSDB; AAQ24681.
 XX PT Compsns. contg. polypeptide antigen recognised by antibody H23 - for
 PT treatment of mammary tumours, also for pox virus compns. for use in
 PT vaccines.
 XX PS Claim 3; Page 19-21; 29pp; French.
 XX The tumour antigen recognised by antibody H23 is aberrantly expressed in
 CC epithelial cells from cancerous mammary tissue in about 90 per cent of
 CC compared to the amount in the control is associated with the presence of

CC breast cancer cases; in a normal individual expression is negligible. The
 CC antigen exists in two forms: transmembrane (ETA-T) and secreted (ETA-S).
 CC Both forms show a high degree of polymorphism. A 20 amino acid subunit in
 CC ETA can be tandemly repeated up to 80 times. (N.B. two tandem repeats are
 CC shown here; the first half of the amino acid sequence, i.e. on the N-
 CC terminal side of the repeat region, is given in AAR27663). From one
 CC subunit to the next, 1 to 3 amino acids can differ. See also AAQ24678-
 CC Q24681, AAQ29276-7 and AAR23974-R23981. (Updated on 25-MAR-2003 to
 CC correct PN field.)
 XX SQ Sequence 180 AA;
 Query Match 100.0%; Score 49; DB 2; Length 180;
 Best Local Similarity 100.0%; Pred. No. 2;
 Matches 9; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 AC QY 1 STAPPVHN 9
 DB 43 STAPPVHN 51

RESULT 12
 ADI57759
 ID ADI57759 standard; protein; 256 AA.
 XX DT 22-APR-2004 (first entry)
 XX DB Human breast specific protein (BSP) #36.
 XX KW Human; breast specific protein; BSP; metastasis; breast cancer;
 XX AC ADI57759;
 XX KW cytostatic.
 XX OS Homo sapiens.
 XX PN WO2003106648-A2.
 XX PD 24-DEC-2003.
 XX PR 14-JUN-2002; 2002US-0389327P.
 XX PA (DIAD-) DIADEXUS INC.
 XX PI Salceda S, Macina RA, Turner LR, Sun Y, Liu C;
 XX WPI; 2004-082185/08.
 XX DR N-PSDB; ADI57687.

Claim 12; SEQ ID NO 130; 370pp; English.

XX The invention relates to human breast specific nucleic acids (BSNA) and
 CC the breast specific proteins (BSP) they encode. The nucleic acids are
 CC useful for determining the presence of a BSNA in a sample which involves
 CC contacting the sample with a BSNA under conditions in which the BSNA will
 CC selectively hybridise to BSNA in the sample, and detecting the presence
 CC of a BSNA in a sample which involves contacting the sample with a suitable
 CC reagent under conditions in which the reagent will selectively interact
 CC with the BSP, and detecting the interaction of the reagent with a BSP in
 CC the sample. The nucleic acids and proteins are useful for diagnosing or
 CC monitoring the presence and metastases of breast cancer in a patient,
 CC which involves determining an amount of nucleic acid or protein and
 CC comparing the determined amount of nucleic acid or protein in the sample
 CC of the patient to the amount of a breast specific marker in a normal
 CC control, where a difference in the determined amount in the sample
 CC compared to the amount in the control is associated with the presence of

breast cancer. The sequences are useful for treating a patient with breast cancer, involving administering a composition consisting of a BSA or a BSP to a patient, where the administration induces an immune response against the breast cancer cell expressing the BSA or BSP. This sequence represents a human BSP of the invention.

Sequence 256 AA;

Query Match 100.0%; Score 49; DB 8; Length 256;
Best Local Similarity 100.0%; Pred. No. 2.8; Mismatches 0; Indels 0; Gaps 0; Gaps 0;

Qy 1 STAPPVHN 9
Db 130 STAPPVHN 138

RESULT 13

ARR27655
ID ARR27655 standard; protein; 287 AA.

XX
AC AAR27655;
XX
DT 25-MAR-2003 (revised)
DT 06-NOV-1992 (first entry)

DE Secreted form of H23-ETA antigen.

XX
ETA-S; human epithelial antigen; Monoclonal antibody H23; vaccine;
KW malignant tumour; breast cancer; tandem repeat;
XX
Homo sapiens.

FH Key Location/Qualifiers

FT Peptide 1..21

FT FT /label= signal

FT Protein 22..287

FT FT /label= ETA-T

FT Misc-difference 134

FT FT /label= Pro, Ala

FT FT /note= "natural polymorphism"

FT FT Misc-difference 144

FT FT /label= Thr, Asn

FT FT /note= "natural polymorphism"

FT FT Misc-difference 147

FT FT /label= Pro, Ala

FT FT /note= "natural polymorphism"

PN WO9207000-A1.

XX 30-APR-1992.

XX 23-OCT-1991; 91WO-FR000835.

XX 23-OCT-1990; 90FR-00013101.

XX (TRGE) TRANSGENE SA.

XX Chambon P, Kienny MP, Lathe R, Harreveni M;

XX WPI; 1992-167097/20.

N-PSDB; AAQ29277.

PT Compsns, contg. polypeptide antigen recognised by antibody H23 - for treatment of mammary tumours, also for pox virus compsns. for use in vaccines.

XZ Claim 3; Page 19-21; 29pp; French.

CC The tumour antigen recognised by antibody H23 is aberrantly expressed in epithelial cells from cancerous mammary tissue in about 90 per cent of breast cancer cases; in a normal individual expression is negligible. The antigen exists in two forms: transmembrane (ETA-T) and secreted (ETA-S).

CC Both forms show a high degree of polymorphism. A 20 amino acid subunit in ETA can be tandemly repeated up to 80 times. From one subunit to the next, 1 to 3 amino acids can differ. DNA coding for immunogenic fragments of ETA can be inserted into e.g. vaccinia viruses for treatment of mammary tumours. See also AAQ24678-Q24681, AAQ29276-7 and AAK23974- R23981. (Updated on 25-MAR-2003 to correct PN field.)

SQ Sequence 287 AA;

Query Match 100.0%; Score 49; DB 2; Length 287;
Best Local Similarity 100.0%; Pred. No. 3.2; Mismatches 0; Indels 0; Gaps 0;

Qy 1 STAPPVHN 9

Db 150 STAPPVHN 158

RESULT 14

AYY71027
ID AYY71027 standard; protein; 295 AA.

XX
AC AYY71027;
XX
DT 12-SEP-2003 (revised)
DT 29-AUG-2000 (first entry)

XX Ubiquitin-E. coli Laci-human Mucin 1 fusion protein #2.
XX Ubiquitin; Laci; beta-galactosidase; fusion protein; human; Mucin 1;
KW MUC-1; tumour; pMRS30 expression vector; anti-tumour; therapy;
KW immune response; cytostatic; vaccine.

XX Homo sapiens.
OS Escherichia coli.
OS Chimeric.

XX Key Location/Qualifiers
1..123
FT FT /label= UBILaci protein
FT FT /note= "contains ubiquitin sequence fused to a portion of
E. coli Laci"
FT FT Region
124..295
FT FT /label= Human_MUC-1_fragment

XX WO2002025827-A2.

PN WO2002025827-A2.

XX PD 11-MAY-2000.

XX PF 18-OCT-1999; 99WO-EP007874.
XX PD 11-MAY-2000.

XX PR 30-OCT-1998; 98IT-MI02330.

XX PA (MEN) MENARINI RICERCHE SPA.

XX XX
XX PI Parente D, Di Massimo AM, De Santis R;
XX XX
XX DR WPI; 2000-165410/31.
XX N-PSDB; AAD00391.

XX The present sequence is a fusion protein consisting of human Mucin 1 (MUC-1) fragment fused to UBILaci sequence at the N-terminus. The UBILaci sequence consists of ubiquitin from MCF7 cell line and a portion of E. coli beta-galactosidase (lacZ). MUC-1 is an antigenic protein overexpressed in tumour cells. The corresponding DNA sequence is cloned into a pMRS30 expression vector and used in pharmaceutical composition e.g. vaccine for inducing an antigen-specific anti-tumour immune

CC Claim 8; 56pp; English.

CC The present sequence is a fusion protein consisting of UBILaci sequence at the N-terminus. The UBILaci sequence consists of ubiquitin from MCF7 cell line and a portion of E. coli beta-galactosidase (lacZ). MUC-1 is an antigenic protein overexpressed in tumour cells. The corresponding DNA sequence is cloned into a pMRS30 expression vector and used in pharmaceutical composition e.g. vaccine for inducing an antigen-specific anti-tumour immune

CC response. Composition containing this DNA molecule is useful in anti-
 CC tumour therapy of patients affected with tumours characterised by high
 CC MUC-1 expression. (Updated on 12-SEP-2003 to standardise OS field)

XX Sequence 295 AA;

Query Match Score 49; DB 3; Length 295;
 Best Local Similarity 100.0%; Pred. No. 3.3; Mismatches 0; Indels 0; Gaps 0;
 Matches 9; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 Qy 1 STAPPVHN 9
 Db 249 STAPPVHN 257

RESULT 15

ADA50571
 ID ADA50571 standard; protein; 307 AA.

XX ADA50571;

XX

DR 20-NOV-2003 (first entry)

XX

DE Mucin 1 (MUC-1) splice variant #1, SEQ ID NO:26.

XX

Nucleic acid vaccine; DNA vaccine; tumour antigen; Cytokine adjuvant;
 KW humoral response; cellular response; immune response; immunotherapy;
 KW cancer; cytostatic; vaccine; gene therapy; mucin 1; MUC-1.
 XX Unidentified.

OS

XX

PN WO2003031569-A2.

XX

PD 17-APR-2003.

XX

PP 18-SEP-2002; 2002WO-US029640.

XX

PR 10-OCT-2001; 2001US-0328371P.

XX

PA (CENZ) CENTOCOR INC.

XX

Snyder L, Scallion B, Knight DM, McCarthy SG, Goletz TJ;
 Branigan PJ;

XX

DR WPI: 2003-393437/37.

XX

DR N-PSDB; ADA50572.

XX

PT New nucleic acid vaccine, useful for eliciting an immune response to a

PT cancer associated tumor protein in a mammal.

XX

PS Claim 1a; Page 38; 92PP; English.

XX

The invention relates to a nucleic acid vaccine comprising one or more
 CC tumour antigen-encoding nucleic acids and one or more cytokine adjuvant-
 CC encoding nucleic acids. The tumour antigen encoded by the vaccine is
 CC mucin 1 (MUC-1), the kallikrein KLK2, or prostate specific antigen (PSA,
 CC also known as KLU3), and the cytokine adjuvant encoded can be interleukin
 CC -12 (IL-12), granulocyte macrophage-colony stimulating factor (GM-CSF),
 CC or especially interleukin-18. The antigen-encoding nucleic acid
 CC is preferably under the control of a promoter such as the cytomegalovirus
 CC immediate early promoter, the dihydrofolate reductase promoter or the
 CC early or late SV40 promoters. The invention also encompasses the method
 CC of eliciting an immune response to a tumour antigen in a mammal using the
 CC vaccine of the invention. Coexpression of the antigen and adjuvant
 CC induces humoral or cellular response to the tumour antigen, generating
 CC an immune response useful for treatment or prophylaxis of cancers. The
 CC present sequence represents a mucin 1 (MUC-1) polypeptide sequence which
 CC is specifically claimed for use in the vaccine of the invention.

XX Sequence 307 AA;

Query Match Score 49; DB 6; Length 307;
 Best Local Similarity 100.0%; Pred. No. 3.4;

Qy 1 STAPPVHN 9
 Db 170 STAPPVHN 178

Search completed: December 9, 2004, 13:54:22
 Job time : 153 secs

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OM protein - protein search, using sw model

Run on: December 9, 2004, 13:49:19 ; Search time 38 Seconds
(without alignments)

15.707 Million cell updates/sec

Title: US-10-019-513-1

Perfect score: 49

Sequence: 1 STAPPVHN 9

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 478139 seqs, 66318000 residues

Total number of hits satisfying chosen parameters: 478139

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%

Listing first 45 summaries

Database : Issued Patents AA:*

1: /cgn2_6_ptodata/1/iaa/5A_COMB.pep:*

2: /cgn2_6_ptodata/1/iaa/5B_COMB.pep:*

3: /cgn2_6_ptodata/1/iaa/6A_COMB.pep:*

4: /cgn2_6_ptodata/1/iaa/6B_COMB.pep:*

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6: /cgn2_6_ptodata/1/iaa/backfiles.Comb.pep:*

pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query	Match	Length	DB ID	Description
1	49	100.0	1867	2	US-08-479-537A-5	Sequence 5, App1
2	49	100.0	1867	3	US-09-083-116-5	Sequence 5, App1
3	49	100.0	1867	3	US-09-134-916A-5	Sequence 5, App1
4	49	100.0	2035	2	US-08-479-537A-2	Sequence 2, App1
5	49	100.0	2035	3	US-09-083-116-2	Sequence 2, App1
6	49	100.0	2035	3	US-09-134-916A-2	Sequence 2, App1
7	39	79.6	9	1	US-08-787-547-55	Sequence 55, App1
8	39	79.6	9	2	US-08-059-19	Sequence 19, App1
9	39	79.6	9	4	US-09-597-8708-45	Sequence 45, App1
10	39	79.6	9	4	US-09-493-232-1	Sequence 1, App1
11	39	79.6	16	3	US-09-043-731-19	Sequence 19, App1
12	39	79.6	19	1	US-08-091-354-3	Sequence 3, App1
13	39	79.6	19	2	US-08-288-059-9	Sequence 9, App1
14	39	79.6	20	2	US-08-288-059-19	Sequence 1, App1
15	39	79.6	20	2	US-08-288-059-32	Sequence 32, App1
16	39	79.6	20	2	US-08-903-516-20	Sequence 20, App1
17	39	79.6	20	2	US-08-833-807-1	Sequence 1, App1
18	39	79.6	20	3	US-09-333-944-1	Sequence 1, App1
19	39	79.6	20	3	US-08-737-896-3	Sequence 3, App1
20	39	79.6	20	3	US-09-223-043-1	Sequence 1, App1
21	39	79.6	20	3	US-08-134-198E-34	Sequence 34, App1
22	39	79.6	20	4	US-08-847-185-20	Sequence 20, App1
23	39	79.6	20	4	US-09-597-870A-1	Sequence 1, App1
24	39	79.6	20	4	US-09-648-028-40	Sequence 40, App1
25	39	79.6	20	4	US-09-493-232-9	Sequence 9, App1
26	39	79.6	20	4	US-09-493-232-11	Sequence 11, App1
27	39	79.6	20	4	US-09-651-265-1	Sequence 1, App1

ALIGNMENTS

RESULT 1
US-08-479-537A-5
; Sequence 5, Application US/08479537A
; Patent No. 5861381

; GENERAL INFORMATION:
; APPLICANT: CHAMBON, Pierre
; APPLICANT: KIENY, Marie-Paule
; APPLICANT: LATHE, Richard
; APPLICANT: HARVEY, Marc
; TITLE OF INVENTION: PHARMACEUTICAL COMPOSITION FOR THE TREATMENT OR PREVENTION OF A MALIGNANT TUMOR
; NUMBER OF SEQUENCES: 5

; CORRESPONDENCE ADDRESS:
; ADDRESSEE: BURNS, DOANE, SWECKER & MATHIS, L.L.P.
; STREET: P.O. Box 1404
; CITY: Alexandria
; STATE: Virginia
; COUNTRY: United States
; ZIP: 22313-1104

; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent in Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/479,537A
; FILING DATE: 07-JUN-1995
; CLASSIFICATION: 514
; PRIORITY APPLICATION DATA:
; APPLICATION NUMBER: FR 90/13101
; FILING DATE: 23-OCT-1990
; PRIORITY APPLICATION DATA:
; APPLICATION NUMBER: WO PCT/FR91/00835
; FILING DATE: 23-OCT-1991
; PRIORITY APPLICATION DATA:
; APPLICATION NUMBER: US 08/039,320
; FILING DATE: 04-APR-1993
; PRIORITY APPLICATION DATA:
; APPLICATION NUMBER: US 08/403,576
; FILING DATE: 14-MAR-1995
; ATTORNEY/AGENT INFORMATION:
; NAME: Tebkin, Robin L.
; REGISTRATION NUMBER: 15,030
; REFERENCE/DOCKET NUMBER: 017753-025
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (703) 836-6620
; TELEFAX: (703) 836-2021
; SEQUENCE CHARACTERISTICS:
; LENGTH: 1867 amino acids

Page 2

NAME/KEY: Peptide
 LOCATION: 128..1899
 OTHER INFORMATION: /note= "The amino acids spanning 128 to 1899 constitute a repeated region wherein the repeat 20 amino acids, 17 of which are fixed. The number of such repeats varies from 1 to 40."
 FEATURE: Peptide
 NAME/KEY: Peptide
 LOCATION: 134
 OTHER INFORMATION: /note= "Amino acid 134 is X1 = Xaa Xaa Xaa which is the codon for Pro or Ala wherein Pro = CCT, CCC, CCA, or CCG; and Ala = GCT, GCC, GCA, or GCG."
 FEATURE: Peptide
 NAME/KEY: Peptide
 LOCATION: 144
 OTHER INFORMATION: /note= "Amino acid 144 is Y = Xaa which is the codon for Thr or Asn wherein Thr = ACT, ACC, ACA OTHER INFORMATION: or ACG; and Asn = AAT or AAC."
 FEATURE: Peptide
 NAME/KEY: Peptide
 LOCATION: 147
 OTHER INFORMATION: /note= "Amino acid 147 is X2 = Xaa which is the codon for Pro or Ala wherein Pro = CCT, CCC, CCA or CCG; and Ala = GCT, GCC, GCA, or GCG."
 FEATURE: Peptide
 NAME/KEY: Peptide
 LOCATION: 1..21
 OTHER INFORMATION: /note= "Amino acids 1 to 21 are a 21 amino acid precursor sequence."
 US-08-479-537-A2

Query Match 100.0%; Score 49; DB 2; Length 2035;
 Best Local Similarity 100.0%; Pred. No. 5,6.; Mismatches 0; Indels 0; Gaps 0;

Qy 1 STAPPVHN 9
 Db 1730 STAPPVHN 1738

RESULT 5
 US-09-083-116-2
 Sequence 2, Application US/09083116
 Ptent No. 6203195
 GENERAL INFORMATION:
 APPLICANT: CHAMSON, Pierre
 APPLICANT: KIENY, Marie-Paule
 APPLICANT: LATHE, Richard
 APPLICANT: HAREVONI, Mara
 TITLE OF INVENTION: PHARMACEUTICAL COMPOSITION FOR THE TREATMENT OR PREVENTION OF A MALIGNANT TUMOR
 NUMBER OF SEQUENCES: 5
 CORRESPONDENCE ADDRESS:
 ADDRESSEE: BURNS, DOANE, SWECKER & MATHIS, L.L.P.
 STREET: P.O. Box 1404
 CITY: Alexandria
 STATE: Virginia
 COUNTRY: United States
 ZIP: 22313-1404
 COMPUTER READABLE FORM:
 MEDIUM TYPE: Floppy disk
 COMPUTER: IBM PC compatible
 OPERATING SYSTEM: PC-DOS/MS-DOS
 SOFTWARE: PatentIn Release #1.0, Version #1.30
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/09/083,116
 FILING DATE:
 CLASSIFICATION:
 PRIOR APPLICATION DATA:
 APPLICATION NUMBER: 08/479,537
 FILING DATE:
 PRIOR APPLICATION DATA:
 APPLICATION NUMBER: WO PCT/ER91/00835

NAME/KEY: Peptide
 LOCATION: 128..1899
 OTHER INFORMATION: /note= "The amino acids spanning 128 to 1899 constitute a repeated region wherein the repeat 20 amino acids, 17 of which are fixed. The number of such repeats varies from 1 to 40."
 FEATURE: Peptide
 NAME/KEY: Peptide
 LOCATION: 128..1899
 OTHER INFORMATION: /note= "The amino acids spanning 128 to 1899 constitute a repeated region wherein the repeat 20 amino acids, 17 of which are fixed. The number of such repeats varies from 1 to 40."
 FEATURE: Peptide
 NAME/KEY: Peptide
 LOCATION: 134
 OTHER INFORMATION: /note= "Amino acid 134 is X1 = Xaa Xaa Xaa which is the codon for Pro or Ala wherein Pro = CCT, CCC, CCA, or CCG."
 FEATURE: Peptide
 NAME/KEY: Peptide
 LOCATION: 144
 OTHER INFORMATION: /note= "Amino acid 144 is Y = Xaa which is the codon for Thr or Asn wherein Thr = ACT, ACC, ACA or GCG."
 FEATURE: Peptide
 NAME/KEY: Peptide
 LOCATION: 147
 OTHER INFORMATION: /note= "Amino acid 147 is X2 = Xaa which is the codon for Pro or Ala wherein Pro = CCT, CCC, CCA or CCG; and Ala = GCT, GCC, GCA, or GCG."
 FEATURE: Peptide
 NAME/KEY: Peptide
 LOCATION: 147
 OTHER INFORMATION: /note= "Amino acid 147 is X2 = Xaa which is the codon for Thr or Asn wherein Thr = ACT, ACC, ACA or GCG; and Ala = GCT, GCC, GCA, or GCG."
 FEATURE: Peptide
 NAME/KEY: Peptide
 LOCATION: 1..21
 OTHER INFORMATION: /note= "Amino acids 1 to 21 are a 21 amino acid precursor sequence."
 US-09-083-116-2

Query Match 100.0%; Score 49; DB 3; Length 2035;
 Best Local Similarity 100.0%; Pred. No. 5,6.; Mismatches 0; Indels 0; Gaps 0;

Qy 1 STAPPVHN 9
 Db 1730 STAPPVHN 1738

RESULT 6
 US-09-134-916A-2
 Sequence 2, Application US/09134916A
 Patent No. 6328956
 GENERAL INFORMATION:
 APPLICANT: CHAMON, Pierre
 APPLICANT: KIENY, Marie-Paule
 APPLICANT: LATHE, Richard
 APPLICANT: HAREVONI, Mara
 TITLE OF INVENTION: PHARMACEUTICAL COMPOSITION FOR THE TREATMENT OR PREVENTION OF A MALIGNANT TUMOR
 TITLE OF INVENTION: TREATMENT OR PREVENTION OF A MALIGNANT TUMOR

APPLICANT: PINN, OLIVERA J.
 APPLICANT: FONTEENOT, J. D.
 APPLICANT: MONTELARO, RONALD C.
 TITLE OF INVENTION: SYNTHETIC MULTIPLE TANDEM REPEAT MUCIN
 TITLE OF INVENTION: AND MUCIN-LIKE PEPTIDES, AND USES THEREOF
 NUMBER OF SEQUENCES: 36
 CORRESPONDENCE ADDRESS:
 ADDRESSEE: CUSHMAN DARBY & CUSHMAN, L.L.P.
 STREET: 1100 NEW YORK AVENUE, N.W.
 CITY: WASHINGTON
 STATE: D.C.
 COUNTRY: USA
 ZIP: 20005
 COMPUTER READABLE FORM:
 MEDIUM TYPE: FLOPPY DISK
 COMPUTER: IBM PC compatible
 OPERATING SYSTEM: PC-DOS/MS-DOS
 SOFTWARE: PatentIn Release #1.0, Version #1.25
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/08/288,059
 FILING DATE: 08-AUG-1994
 CLASSIFICATION: 424
 ATTORNEY/AGENT INFORMATION:
 NAME: CHAPIN, MARLANA K.
 REGISTRATION NUMBER: 35,843
 REFERENCE/DOCKET NUMBER: 61137/205204
 TELECOMMUNICATION INFORMATION:
 TELEPHONE: 202-861-3711
 TELEFAX: 202-22-0344
 TELEX: 6714627 CUSH
 INFORMATION FOR SEQ ID NO: 19:
 SEQUENCE CHARACTERISTICS:
 LENGTH: 9 amino acids
 TYPE: amino acid
 STRANDEDNESS: Single
 TOPOLOGY: Linear
 MOLECULE TYPE: Peptide
 US-08-288-059-19

Query Match 79.6%; Score 39; DB 2; Length 9;
 Best Local Similarity 77.8%; Pred. No. 3.8e+05; Mismatches 2; Indels 0; Gaps 0;

Qy 1 STAPPVHNV 9
 Db 1 STAPPVHGV 9

RESULT 9
 US-09-593-870A-45
 Sequence 45, Application US/09593870A
 GENERAL INFORMATION:
 APPLICANT: McKenzie, Ian F.C.
 APPLICANT: Apostolopoulos, Vassilios
 APPLICANT: Pietersz, Geoff Allan
 TITLE OF INVENTION: Antigen Carbohydrate Compounds and Their
 TITLE OF INVENTION: Use in Immunotherapy
 FILE REFERENCE: 2368-McKenzie
 CURRENT APPLICATION NUMBER: US/09/593,870A
 CURRENT FILING DATE: 2000-16-14
 PRIOR APPLICATION NUMBER: 09/223,043
 PRIOR FILING DATE: 1998-12-30
 NUMBER OF SEQ ID NOS: 69
 SOFTWARE: FastSEQ for Windows Version 3.0
 SEQ ID NO: 45
 LENGTH: 9
 TYPE: PRT
 ORGANISM: Homo sapiens

Query Match 79.6%; Score 39; DB 4; Length 9;
 Best Local Similarity 77.8%; Pred. No. 3.8e+05;

Qy 1 STAPPVHNV 9
 Db 1 STAPPVHGV 9

RESULT 11
 US-09-04-731-19
 Sequence 19, Application US/09043731A
 GENERAL INFORMATION:
 APPLICANT: The Austin Research Institute
 TITLE OF INVENTION: Mimicking Peptides in Cancer Therapy

APPLICANT: AGRAWAL, Babita
 KRANTZ, Mark J.
 REDDISH, Mark A.
 LONGENECKER, B. Michael
 TITLE OF INVENTION: METHOD FOR GENERATING ACTIVATED T-CELLS
 AND ANTIGEN-PULSED ANTIGEN-PRESENTING CELLS
 NUMBER OF SEQUENCES: 34
 CORRESPONDENCE ADDRESS:
 ADDRESSEE: FOLEY & LARDNER
 STREET: 3000 K Street, N.W.
 CITY: Washington
 STATE: D.C.
 COUNTRY: U.S.A.
 ZIP: 20007-5109
 COMPUTER READABLE FORM:
 MEDIUM TYPE: Floppy disk
 COMPUTER: IBM PC compatible
 OPERATING SYSTEM: PC-DOS/MS-DOS
 SOFTWARE: PatentIn Release #1.0, Version #1.30
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/09/497,232
 FILING DATE: 03-Feb-2000
 CLASSIFICATION: <Unknown>
 PRIOR APPLICATION DATA:
 APPLICATION NUMBER: US/09/074,410
 FILING DATE: 03-MAY-1998
 APPLICATION NUMBER: US/09/497,232
 FILING DATE: 03-MAY-1997
 ATTORNEY/AGENT INFORMATION:
 NAME: Saxe, Bernhard D.
 REGISTRATION NUMBER: 28,665
 REFERENCE/DOCKET NUMBER: 042881/0114
 TELECOMMUNICATION INFORMATION:
 TELEPHONE: (202) 672-5300
 TELEFAX: (202) 672-5399
 INFORMATION FOR SEQ ID NO: 1:
 SEQUENCE CHARACTERISTICS:
 LENGTH: 9 amino acids
 TYPE: amino acid
 STRANDEDNESS: <Unknown>
 TOPOLOGY: Linear
 MOLECULE TYPE: peptide
 SEQUENCE DESCRIPTION: SEQ ID NO: 1:
 US-09-497-232-1

Query Match 79.6%; Score 39; DB 4; Length 9;
 Best Local Similarity 77.8%; Pred. No. 3.8e+05; Mismatches 2; Indels 0; Gaps 0;

Qy 1 STAPPVHNV 9
 Db 1 STAPPVHGV 9

FILE REFERENCE: CALA-200
 CURRENT APPLICATION NUMBER: US/09/043,731A
 NUMBER OF SEQ ID NOS: 26
 SOFTWARE: PatentIn Ver. 2.0
 SEQ ID NO: 19
 LENGTH: 16
 TYPE: PRT
 ORGANISM: Artificial Sequence
 FEATURE: Description of Artificial Sequence: single
 OTHER INFORMATION: Stranded linear peptide
 US-09-043-731-19

Query Match Score 79.6%; DB 3; Length 16;
 Best Local Similarity 77.8%; Prod. No. 1.7%;
 Matches 7; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
 Query 1 STAPPVINY 9
 Db 5 STAPPAHGV 13

RESULT 12
 US-08-099-354-3
 Sequence 3, Application US/08099354
 Patent No. 5744144
 GENERAL INFORMATION:
 APPLICANT: FINN, OLIVERA J.
 APPLICANT: FONTENOT, J. D.
 APPLICANT: MONTELARO, RONALD C.
 TITLE OF INVENTION: SYNTHETIC MULTIPLE TANDEM REPEAT MUCIN
 NUMBER OF SEQUENCES: 10
 TITLE OF INVENTION: SYNTHETIC MULTIPLE TAMDEM REPEAT MUCIN
 NUMBER OF SEQUENCES: 35
 TITLE OF INVENTION: AND MUCIN-LIKE PEPTIDES, AND USES THEREOF
 NUMBER OF SEQUENCES: 10
 ADDRESS: CUSHMAN, DARBY & CUSHMAN
 STREET: 1100 NEW YORK AVENUE, N.W.
 CITY: WASHINGTON
 STATE: D.C.
 COUNTRY: USA
 ZIP: 20005

COMPUTER READABLE FORM:
 MEDIUM TYPE: Floppy disk
 COMPUTER: IBM PC compatible
 OPERATING SYSTEM: PC-DOS/MS-DOS
 SOFTWARE: PatentIn Release #1.0, Version #1.25
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/08/288,059
 FILING DATE: 08-AUG-1994
 CLASSIFICATION: 424
 ATTORNEY/AGENT INFORMATION:
 NAME: CHAPIN, MARILIA K.
 REGISTRATION NUMBER: 35,843
 REFERENCE/DOCKET NUMBER: 61137/205204
 TELECOMMUNICATION INFORMATION:
 TELEPHONE: 202-861-3711
 TELEFAX: 202-822-0944
 TELEX: 6714627 CUSH
 INFORMATION FOR SEQ ID NO: 9:
 SEQUENCE CHARACTERISTICS:
 LENGTH: 19 amino acids
 TYPE: amino acid
 STRANDEDNESS: single
 TOPOLOGY: linear
 MOLECULE TYPE: peptide
 US-08-288-059-9

Query Match Score 79.6%; DB 2; Length 19;
 Best Local Similarity 77.8%; Prod. No. 2; Mismatches 0; Indels 2; Gaps 0;
 Query 1 STAPPVINY 9
 Db 8 STAPPAHGV 16

RESULT 13
 US-08-288-059-1
 Sequence 9, Application US/08288059
 Patent No. 5827666
 GENERAL INFORMATION:
 APPLICANT: FINN, OLIVERA J.
 APPLICANT: FONTENOT, J. D.
 APPLICANT: MONTELARO, RONALD C.
 TITLE OF INVENTION: SYNTHETIC MULTIPLE TANDEM REPEAT MUCIN
 NUMBER OF SEQUENCES: 36
 TITLE OF INVENTION: AND MUCIN-LIKE PEPTIDES, AND USES THEREOF
 ADDRESS: CUSHMAN, DARBY & CUSHMAN, L.L.P.
 STREET: 1100 NEW YORK AVENUE, N.W.
 CITY: WASHINGTON
 STATE: D.C.
 COUNTRY: USA

Query Match Score 79.6%; DB 1; Length 19;
 Best Local Similarity 77.8%; Prod. No. 2; Mismatches 0; Indels 0; Gaps 0;
 Query 1 STAPPVINY 9

ZIP: 20005
 COMPUTER READABLE FORM:
 MEDIUM TYPE: Floppy disk.
 COMPUTER: IBM PC compatible
 OPERATING SYSTEM: PC-DOS/MS-DOS
 SOFTWARE: PatentIn Release #1.0, Version #1.25
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/08/288,059
 FILING DATE: 08-AUG-1994
 CLASSIFICATION: 424
 ATTORNEY/AGENT INFORMATION:
 NAME: CHAPIN, MARLANA K.
 REGISTRATION NUMBER: 35,843
 REFERENCE/DOCKET NUMBER: 61137/205204
 TELECOMMUNICATION INFORMATION:
 TELEPHONE: 202-861-4711
 TELEX: 671462Z CUSH
 INFORMATION FOR SEQ ID NO: 1:
 SEQUENCE CHARACTERISTICS:
 LENGTH: 20 amino acids
 TYPE: amino acid
 STRANDEDNESS: single
 TOPOLOGY: linear
 MOLECULE TYPE: Peptide
 US-08-288-059-1

Query Match 79.6%; Score 39; DB 2; Length 20;
 Best Local Similarity 77.8%; Pred. No. 2.1;
 Matches 7; Conservative 0; Mismatches 2;
 Indels 0; Gaps 0;

Qy 1 STAPPVNV 9
 Db 9 STAPPVNV 17

RESULT 15
 US-08-288-059-32
 Sequence 32, Application US/08288059
 Patent No. 5,827,666
 GENERAL INFORMATION:
 APPLICANT: FINN, OLIVERA J.
 APPLICANT: FONTENOT, J. D.
 APPLICANT: MONTEJARO, RONALD C.
 TITLE OF INVENTION: SYNTHETIC MULTIPLE TANDEM REPEAT MUCIN
 NUMBER OF SEQUENCES: 36
 CORRESPONDENCE ADDRESS:
 ADDRESSEE: CUSHMAN, DARBY & CUSHMAN, L.L.P.
 STREET: 1100 NEW YORK AVENUE, N.W.
 CITY: WASHINGTON
 STATE: D.C.
 COUNTRY: USA
 ZIP: 20005
 COMPUTER READABLE FORM:
 MEDIUM TYPE: Floppy disk.
 COMPUTER: IBM PC compatible
 OPERATING SYSTEM: PC-DOS/MS-DOS
 SOFTWARE: PatentIn Release #1.0, Version #1.25
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/08/288,059
 FILING DATE: 08-AUG-1994;
 CLASSIFICATION: 424
 ATTORNEY/AGENT INFORMATION:
 NAME: CHAPIN, MARLANA K.
 REGISTRATION NUMBER: 35,843
 REFERENCE/DOCKET NUMBER: 61137/205204
 TELECOMMUNICATION INFORMATION:
 TELEPHONE: 202-861-3711
 TELEX: 671462Z CUSH
 INFORMATION FOR SEQ ID NO: 32:
 SEQUENCE CHARACTERISTICS:

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OM protein - protein search, using sw model

Run on: December 9, 2004, 13:57:50 ; Search time 144 Seconds
(without alignments)

22.324 Million cell updates/sec

Title: US-10-019-513-1

Perfect score: 49

Sequence: 1 STAPPVHN 9

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 1585576 seqs, 35178320 residues

Total number of hits satisfying chosen parameters: 1585576

Minimum DB seq length: 0

Maximum DB seq length: 20000000000

Post-processing: Minimum Match 0% Maximum Match 100%

Listing first 45 summaries

Published Applications AA:*

1: /cgn2_6/ptodata/2/pubpaas/US07_PUBCOMB.pep:*

2: /cgn2_6/ptodata/2/pubpaas/US07_NEW_PUB.pep:*

3: /cgn2_6/ptodata/2/pubpaas/US06_PUB.pep:*

4: /cgn2_6/ptodata/2/pubpaas/US06_PUBCOMB.pep:*

5: /cgn2_6/ptodata/2/pubpaas/US08_PUB.pep:*

6: /cgn2_6/ptodata/2/pubpaas/US08_PUBCOMB.pep:*

7: /cgn2_6/ptodata/2/pubpaas/US10_PUB.pep:*

8: /cgn2_6/ptodata/2/pubpaas/US10_PUBCOMB.pep:*

9: /cgn2_6/ptodata/2/pubpaas/US09A_PUBCOMB.pep:*

10: /cgn2_6/ptodata/2/pubpaas/US09B_PUBCOMB.pep:*

11: /cgn2_6/ptodata/2/pubpaas/US09C_PUBCOMB.pep:*

12: /cgn2_6/ptodata/2/pubpaas/US09_NEW_PUB.pep:*

13: /cgn2_6/ptodata/2/pubpaas/US10A_PUBCOMB.pep:*

14: /cgn2_6/ptodata/2/pubpaas/US10B_PUBCOMB.pep:*

15: /cgn2_6/ptodata/2/pubpaas/US10C_PUBCOMB.pep:*

16: /cgn2_6/ptodata/2/pubpaas/US10D_PUBCOMB.pep:*

17: /cgn2_6/ptodata/2/pubpaas/US10_NEW_PUB.pep:*

18: /cgn2_6/ptodata/2/pubpaas/US11_NEW_PUB.pep:*

19: /cgn2_6/ptodata/2/pubpaas/US60_NEW_PUB.pep:*

20: /cgn2_6/ptodata/2/pubpaas/US60_PUBCOMB.pep:*

Sequence 10, Appl

Sequence 15, Appl

Sequence 311, Appl

Sequence 120, Appl

Sequence 1210, Appl

Sequence 851, Appl

Sequence 267125, Appl

Sequence 55, Appl

Sequence 55, Appl

Sequence 45, Appl

Sequence 20, Appl

Sequence 1, Appl

Sequence 11, Appl

Sequence 1, Appl

Sequence 1, Appl

Sequence 44, Appl

Sequence 47, Appl

Sequence 138, Appl

Sequence 20, Appl

Sequence 1, Appl

Sequence 97, Appl

Sequence 44, Appl

Sequence 47, Appl

Sequence 17, Appl

Sequence 20, Appl

Sequence 40, Appl

Sequence 31, Appl

Sequence 3, Appl

Sequence 31, Appl

Sequence 215, Appl

Sequence 32, Appl

Sequence 196, Appl

Sequence 207, Appl

Sequence 1, Appl

Sequence 1, Appl

Sequence 24, Appl

ALIGNMENTS

RESULT 1
US-10-247-703-43

/ Sequence 43, Application US/10247-703-43

/ Publication No. US20030635924

/ GENERAL INFORMATION:

/ APPLICANT: Branigan, Peter J

/ APPLICANT: Goletz, Teresa J

/ APPLICANT: Knight, David M

/ APPLICANT: McCarthy, Stephen G

/ APPLICANT: Schilder, Bernard J

/ APPLICANT: Snyder, Linda A

/ TITLE OF INVENTION: NUCLEIC ACID VACCINES USING TUMOR ANTIGEN ENCODING NUCLEIC ACIDS

/ TITLE OF INVENTION: CYTOKINE ADJUVANT ENCODING NUCLEIC ACID

/ FILE REFERENCE: CEN310

/ CURRENT APPLICATION NUMBER: US/10/247,703

/ CURRENT FILING DATE: 2002-09-20

/ PRIOR APPLICATION NUMBER: 60/328,371

/ PRIOR FILING DATE: 2001-10-10

/ NUMBER OF SEQ ID NOS: 77

Result No.	Score	Query Match	Length	DB ID	Description
1	49	100.0	9	14 US-10-247-703-43	Sequence 43, Appl
2	49	100.0	9	15 US-10-47-161-98	Sequence 98, Appl
3	49	100.0	13	14 US-10-07-136-18	Sequence 18, Appl
4	49	100.0	30	15 US-10-26-734-1168	Sequence 1168, Appl
5	49	100.0	307	14 US-10-24-703-26	Sequence 26, Appl
6	49	100.0	312	15 US-10-26-734-824	Sequence 824, Appl
7	49	100.0	475	14 US-10-24-703-22	Sequence 22, Appl
8	49	100.0	475	14 US-10-47-312-1	Sequence 1, Appl
9	49	100.0	508	14 US-10-07-136-20	Sequence 20, Appl
10	49	100.0	515	14 US-10-247-703-20	Sequence 21, Appl
11	49	100.0	515	14 US-10-09-340-212	Sequence 156, Appl
12	49	100.0	515	14 US-10-171-311-156	Sequence 19, Appl
13	49	100.0	515	15 US-10-612-090-19	

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Qy	Score	Match	Length	DB ID	Description
1	49	100.0	9	14 US-10-247-703-43	Sequence 43, Appl
2	49	100.0	9	15 US-10-47-161-98	Sequence 98, Appl

Query Match 100.0% ; Score 49; DB 14; Length 9;

Best Local Similarity 10.0% ; Pred. No. 1.4e+06; Mismatches 0; Indels 0; Gaps 0;

OS-10-247-703-43

OS/10247-703-43

OS/20030635924

OS/10-247-703-43

OS/10-07-136-18

OS/10-26-734-1168

OS/10-24-703-26

OS/10-26-734-824

OS/10-24-703-22

OS/10-47-312-1

OS/10-07-136-20

OS/10-247-703-20

OS/10-09-340-212

OS/10-171-311-156

OS/10-612-090-19

STAPPVHN 9

STAPPVHN 9

RESULT 2

US-10-447-161-98
; Sequence 98, Application US/10447161
; Publication No. US20040023314A1
; GENERAL INFORMATION:
; APPLICANT: Wang, Rong-fu
; TITLE OF INVENTION: Mutant Fibronectin and Tumor Metastasis
; FILE REFERENCE: HO-P0248AUS1
; CURRENT APPLICATION NUMBER: US/10/447.161
; CURRENT FILING DATE: 2003-05-28
; PRIORITY NUMBER: 60/383,530
; PRIORITY FILING DATE: 2002-05-28
; NUMBER OF SEQ ID NOS: 148
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO: 98
; LENGTH: 9
; TYPE: PRT
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Synthetic Peptide

US-10-447-161-98

Query Match 100.0%; Score 49; DB 15; Length 9;
Best Local Similarity 100.0%; Pred. No. 1.4e+06;
Matches 9; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 STAPPVHN 9
Db 1 STAPPVHN 9

RESULT 5
US-10-247-703-26
; Sequence 26, Application US/10247703
; Publication No. US2003063597A1
; GENERAL INFORMATION:
; APPLICANT: Branigan, Patrick
; APPLICANT: Goleitz, Theresa J
; APPLICANT: Knight, David M
; APPLICANT: McCarthy, Stephen G
; APPLICANT: Scallion, Bernard J
; APPLICANT: Snyder, Linda A
; TITLE OF INVENTION: NUCLEIC ACID VACCINES USING TUMOR ANTIGEN ENCODING NUCLEIC ACID
; FILE REFERENCE: CEN310
; CURRENT APPLICATION NUMBER: US/10/247.703
; CURRENT FILING DATE: 2002-09-20
; PRIORITY NUMBER: 60/328,371
; PRIORITY FILING DATE: 2001-10-10
; NUMBER OF SEQ ID NOS: 77
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO: 26
; LENGTH: 307
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-247-703-26

Query Match 100.0%; Score 49; DB 14; Length 307;
Best Local Similarity 100.0%; Pred. No. 2.1e+00;
Matches 9; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 STAPPVHN 9
Db 170 STAPPVHN 178

RESULT 6
US-10-296-734-924
; Sequence 824, Application US/10296734
; Publication No. US2004005413A1
; GENERAL INFORMATION:
; APPLICANT: Thompson, Scott A
; APPLICANT: Ramsdaw, Ian A
; TITLE OF INVENTION: Synthetic molecules and uses therefor
; FILE REFERENCE: Savine
; CURRENT APPLICATION NUMBER: US/10/296,734
; CURRENT FILING DATE: 2003-08-04

US-10-296-734-1168
; Sequence 1168, Application US/10296734
; Publication No. US2004005413A1
; GENERAL INFORMATION:

PRIOR APPLICATION NUMBER: AU P07761/00
 PRIOR DATE: 2000-05-26
 NUMBER OF SEQ ID NOS: 1507
 SOFTWARE: PatentIn version 3.2
 SEQ ID NO: 824
 LENGTH: 312
 TYPE: PRT
 ORGANISM: Artificial
 FEATURE:
 OTHER INFORMATION: MUC1R consensus polypeptide
 US-10-296-734-824

Query Match 100.0%; Score 49; DB 15; Length 312;

Best Local Similarity 100.0%; Pred. No. 2,1; Matches 9; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 STAPPVNV 9
 Db 7 STAPPVNV 15

RESULT 7

US-10-247-703-22
 Sequence 22, Application US/10247703
 Publication No. US200300663597A1
 GENERAL INFORMATION:
 APPLICANT: Branigan, Patrick
 APPLICANT: Goletz, Theresa J
 APPLICANT: Knight, David M
 APPLICANT: McCarthy, Stephen G
 APPLICANT: Scallion, Bernard J
 APPLICANT: Snyder, Linda A
 TITLE OF INVENTION: NUCLEAR ACID VACCINES USING TUMOR ANTIGEN ENCODING NUCLEIC ACIDS
 FILE REFERENCE: CEN310
 CURRENT APPLICATION NUMBER: US/10/247,703
 CURRENT FILING DATE: 2002-09-20
 PRIOR APPLICATION NUMBER: 60/328,371
 PRIOR FILING DATE: 2001-10-10
 NUMBER OF SEQ ID NOS: 77
 SOFTWARE: PatentIn version 3.1
 SEQ ID NO: 22
 LENGTH: 475
 TYPE: PRT
 ORGANISM: Homo sapiens
 US-10-247-703-22

Query Match 100.0%; Score 49; DB 14; Length 475;

Best Local Similarity 100.0%; Pred. No. 3,2; Matches 9; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 STAPPVNV 9
 Db 170 STAPPVNV 178

RESULT 8

US-10-417-312-1
 Sequence 1, Application US/10417312
 Publication No. US20030235868A1
 GENERAL INFORMATION:
 APPLICANT: Dyax Corp
 TITLE OF INVENTION: Antibodies Specific for Mucin Polypeptide
 FILE REFERENCE: 2403/2002
 CURRENT APPLICATION NUMBER: US/10/417,312
 CURRENT FILING DATE: 2003-04-16
 PRIOR APPLICATION NUMBER: US 60/374,432
 PRIOR FILING DATE: 2002-04-22
 NUMBER OF SEQ ID NOS: 9
 SOFTWARE: PatentIn version 3.2
 SEQ ID NO: 1
 LENGTH: 475
 TYPE: PRT

; ORGANISM: Homo sapiens
 US-10-417-312-1

Query Match 100.0%; Score 49; DB 14; Length 475;
 Best Local Similarity 100.0%; Pred. No. 3,2;
 Matches 9; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 Qy 1 STAPPVNV 9
 Db 170 STAPPVNV 178

RESULT 9

US-10-057-136-20
 Sequence 20, Application US/10057136
 Publication No. US2003021770A1
 GENERAL INFORMATION:
 APPLICANT: SCHLOM, JEFFREY
 APPLICANT: KANTOR, JUDITH
 APPLICANT: KOFIS, DONALD
 APPLICANT: PANICALI, DENNIS
 APPLICANT: GRITZ, LINDA
 TITLE OF INVENTION: RECOMBINANT POX VIRUS FOR IMMUNIZATION AGAINST MUC1
 TITLE OF INVENTION: TUMOR-ASSOCIATED ANTIGEN
 FILE REFERENCE: 700953/47113C
 CURRENT APPLICATION NUMBER: US/10/057,136
 CURRENT FILING DATE: 2002-01-25
 PRIOR APPLICATION NUMBER: 09/366,670
 PRIOR FILING DATE: 1999-08-03
 PRIOR APPLICATION NUMBER: PCT/US98/03693
 PRIOR FILING DATE: 1998-02-24
 PRIOR APPLICATION NUMBER: 60/038,253
 PRIOR FILING DATE: 1997-02-24
 NUMBER OF SEQ ID NOS: 20
 SEQ ID NO: 20
 LENGTH: 508
 TYPE: PRT
 ORGANISM: Homo sapiens
 US-10-057-136-20

Query Match 100.0%; Score 49; DB 14; Length 508;
 Best Local Similarity 100.0%; Pred. No. 3,4;
 Matches 9; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 Qy 1 STAPPVNV 9
 Db 203 STAPPVNV 211

RESULT 10

US-10-247-703-20
 Sequence 20, Application US/10247703
 Publication No. US20030063597A1
 GENERAL INFORMATION:
 APPLICANT: Branigan, Patrick
 APPLICANT: Goletz, Theresa J
 APPLICANT: Knight, David M
 APPLICANT: McCarthy, Stephen G
 APPLICANT: Scallion, Bernard J
 APPLICANT: Snyder, Linda A
 TITLE OF INVENTION: NUCLEAR ACID VACCINES USING TUMOR ANTIGEN ENCODING NUCLEIC ACID
 FILE REFERENCE: CEN310
 CURRENT APPLICATION NUMBER: US/10/247,703
 CURRENT FILING DATE: 2002-09-20
 PRIOR APPLICATION NUMBER: 60/328,371
 PRIOR FILING DATE: 2001-10-10
 NUMBER OF SEQ ID NOS: 77
 SEQ ID NO: 20
 LENGTH: 515
 TYPE: PRT

ORGANISM: Homo sapiens
US-10-247-703-20

Query Match Score 49; DB 14; Length 515;
Best Local Similarity 100.0%; Pred. No. 3.5%;
Matches 9; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 STAPPVHN 9
Db 210 STAPPVHN 218

RESULT 11
US-10-097-340-212
Sequence 212, Application US/10097340
Publication No. US20030087250A1

GENERAL INFORMATION:
APPLICANT: John MONAHAN

APPLICANT: Manjula GANNAVARAPU
APPLICANT: Sebastian HOERSCH
APPLICANT: Shubhangi KAMATKAR
APPLICANT: Steve G. KOVATS
APPLICANT: Rachel E. MEYERS
APPLICANT: Michael J. MORRISEY
APPLICANT: Peter OLANDT
APPLICANT: Ami SEN
APPLICANT: Peter VIBY

APPLICANT: Gordon B. MILLS
APPLICANT: Robert C. BAST, JR.
APPLICANT: Karen LU
APPLICANT: Rosemarie SCHMANDT
APPLICANT: Xumei ZHAO
APPLICANT: Karen GIATTI

TITLE OF INVENTION: Nucleic Acid Molecules and Proteins For The Identification, Prevention, and Therapy of Ovarian Cancer
FILE REFERENCE: MRI-030

CURRENT APPLICATION NUMBER: US/10/097,340
CURRENT FILING DATE: 2002-03-14
PRIOR APPLICATION NUMBER: 60/276,025
PRIOR FILING DATE: 2001-03-14
PRIOR APPLICATION NUMBER: 60/325,149
PRIOR FILING DATE: 2001-09-26
PRIOR APPLICATION NUMBER: 60/276,026
PRIOR FILING DATE: 2001-03-14
PRIOR APPLICATION NUMBER: 60/324,967
PRIOR FILING DATE: 2001/09/26
PRIOR APPLICATION NUMBER: 60/311,732
PRIOR FILING DATE: 2001-08-10
PRIOR APPLICATION NUMBER: 60/325,102
PRIOR FILING DATE: 2001-09-26
PRIOR APPLICATION NUMBER: 60/323,580
PRIOR FILING DATE: 2001-09-19
SOFTWARE: Fast-SEQ for Windows Version 4.0
SEQ ID NO: 212
TYPE: PRT
ORGANISM: Homo sapiens
US-10-097-340-212

GENERAL INFORMATION:
APPLICANT: Schlegel, Robert
APPLICANT: Chen, Yan
APPLICANT: Zhao, Xumei
APPLICANT: Monahan, John
APPLICANT: Kamatkar, Shubhangi
APPLICANT: Glatt, Karen
APPLICANT: Gannavarapu, Manjula
APPLICANT: Hoersch, Sebastian
APPLICANT: GANNAVARAPU, COMPOSITIONS, KITS, AND METHODS FOR IDENTIFICATION, ASSESSMENT, PREVENTION, AND THERAPY OF CERVICAL CANCER
FILE REFERENCE: MRI-035

CURRENT APPLICATION NUMBER: US/10/171,311
CURRENT FILING DATE: 2002-06-12
PRIOR APPLICATION NUMBER: US 60/298,159
PRIOR FILING DATE: 2001-06-13
PRIOR APPLICATION NUMBER: US 60/298,155
PRIOR FILING DATE: 2001-06-13
PRIOR APPLICATION NUMBER: US 60/335,936
PRIOR FILING DATE: 2001-11-14
NUMBER OF SEQ ID NOS: 238
SOFTWARE: Fast-SEQ for Windows Version 4.0
SEQ ID NO: 156
LENGTH: 515
TYPE: PRT
ORGANISM: Homo sapiens
US-10-171-311-156

Query Match Score 100.0%; Pred. No. 3.5%; Length 515;
Best Local Similarity 100.0%; Mismatches 0; Indels 0; Gaps 0;

Qy 1 STAPPVHN 9
Db 210 STAPPVHN 218

RESULT 13
US-10-612-090-19
Sequence 19, Application US/10/612090
Publication No. US20040057952A1

GENERAL INFORMATION:
APPLICANT: Immunogen, Inc.
TITLE OF INVENTION: ANTIBODIES TO NON-SHEDED MUC1 AND MUC16, AND USES THEREOF
FILE REFERENCE: AB340

CURRENT APPLICATION NUMBER: US/10/612,090
CURRENT FILING DATE: 2003-07-03
PRIOR APPLICATION NUMBER: US 60/393,094
PRIOR FILING DATE: 2002-07-03
NUMBER OF SEQ ID NOS: 33
SOFTWARE: PatentIn version 3.2
SEQ ID NO: 19
LENGTH: 515
TYPE: PRT
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: Exemplary Muc1 protein
US-10-612-090-19

Query Match Score 100.0%; Pred. No. 3.5%; Length 515;
Best Local Similarity 100.0%; Mismatches 0; Indels 0; Gaps 0;

Qy 1 STAPPVHN 9
Db 210 STAPPVHN 218

RESULT 14
US-09-996-069-10
Sequence 10, Application US/09996069
Publication No. US20030036199A1

Query Match Score 100.0%; Pred. No. 3.5%; Length 515;
Best Local Similarity 100.0%; Mismatches 0; Indels 0; Gaps 0;

Qy 1 STAPPVHN 9
Db 210 STAPPVHN 218

GENERAL INFORMATION:
 APPLICANT: Bandid, Cynthia
 APPLICANT: Bandid, R. Shoshana
 TITLE OF INVENTION: DIAGNOSTIC TUMOR MARKERS, DRUG SCREENING FOR TUMORIGENESIS INHIBITORS, AND COMPOSITIONS AND METHODS FOR TREATMENT OF CANCER
 FILE REFERENCE: M0105/70071
 CURRENT APPLICATION NUMBER: US/09/996,069
 CURRENT FILING DATE: 2001-11-27
 NUMBER OF SEQ ID NOS: 35
 SOFTWARE: PatentIn version 3.1
 SEQ ID NO 10
 LENGTH: 1255
 TYPE: PRT
 ORGANISM: Homo sapiens
 US-09-996-069-10

Query Match 100.0%; Score 49; DB 10; Length 1255;
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RESULT 15
 US-10-171-311-158
 Sequence 158, Application US/10171311
 Publication No. US20030087270A1
 GENERAL INFORMATION:
 APPLICANT: Schleger, Robert
 APPLICANT: Chen, Yan
 APPLICANT: Zhao, Xumei
 APPLICANT: Monahan, John
 APPLICANT: Kamatkar, Shubhangi
 APPLICANT: Glatt, Karen
 APPLICANT: Gannavarapu, Manjula
 APPLICANT: Hoersh, Sebastian
 TITLE OF INVENTION: NOVEL GENES, COMPOSITIONS, KITS, AND METHODS FOR IDENTIFICATION, ASSESSMENT, PREVENTION, AND THERAPY OF CERVICAL CANCER
 FILE REFERENCE: M01-035
 CURRENT APPLICATION NUMBER: US/10/171-311
 CURRENT FILING DATE: 2002-06-12
 PRIOR APPLICATION NUMBER: US 60/238,159
 PRIOR FILING DATE: 2001-06-13
 PRIOR APPLICATION NUMBER: US 60/288,155
 PRIOR FILING DATE: 2001-06-13
 PRIOR APPLICATION NUMBER: US 60/335,936
 PRIOR FILING DATE: 2001-11-14
 NUMBER OF SEQ ID NOS: 238
 SOFTWARE: Fast-SEQ for Windows Version 4.0
 SEQ ID NO 158
 LENGTH: 1255
 TYPE: PRT
 ORGANISM: Homo sapiens
 US-10-171-311-158

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Search completed: December 9, 2004, 14:10:02
 Job time : 144 secs

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